Effects of La substitution on superconducting state of CeCoIn₅

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We report effects of La substitution on superconducting state of heavy fermion superconductor CeCoIn₅, as seen in transport and magnetization measurements. As opposed to the case of conventional superconductors, pair breaking by nonmagnetic La results in depression of T_c and indicates strong gap anisotropy. Upper critical field H_{c2} values decrease with increased La concentration, but the critical field anisotropy, $\gamma = H_{c2}^a/H_{c2}^c$, does not change in the Ce_{1-x}La_xCoIn₅ (x=0-0.15). The electronic system is in the clean limit for all values of x.

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I. INTRODUCTION

Heavy fermion superconductors have been extending for more than two decades the affluence of physical phenomena associated with Cooper pair formation.¹ The competition between magnetism and superconductivity for the same Fermi surface of heavily renormalized carriers resulted in observations of unconventional superconductivity^{2,3} and raised speculations that spin pairing might be mediated by magnetic interaction.⁴ Research in the field was associated with difficulties in sample preparation, sample to sample variation, experimental conditions and ultimately, in the number of examples where relevant physical phenomena can be observed in a clean form. The recently discovered CeMIn₅ family (M=Ir, Rh, Co) of heavy fermion superconductors encapsulates many aspects of important physics in this class of materials. CeRhIn₅⁵ superconducts under applied pressures above 17 kbar with T_c around 2 K whereas CeIrIn₅⁶ and CeCoIn₅⁷ are ambient pressure superconductors. CeCoIn₅ offers clean example of ambient pressure heavy fermion superconductivity with a remarkably high $T_c=2.3$ K. The intriguing properties of CeCoIn₅ led to speculation that it may exhibit d-wave superconductivity, 8,9,10 and Fulde-Ferrel-Larkin-Ovchinnikov state in high magnetic fields. ¹¹ In order to have more insight into the nature of CeCoIn₅ we perturbed its superconducting state by substituting La onto the Ce site. For the purpose of comparing influences of magnetic and nonmagnetic pair breaking on T_c suppression, we also substituted 5% of Nd on Ce site. We find that the anisotropy in the upper critical field does not change in the whole concentration range and that the decrease of T_c with increased La doping cannot be explained solely with pressure effects due to unit cell expansion. In addition, our results present an evidence for an anisotropic order parameter in CeCoIn₅.

II. EXPERIMENT

Single crystals of $Ce_{1-x}La_xCoIn_5$ were grown by the self flux method in a manner previously described.⁷ Crys-

tals grew as thin plates with the c axis perpendicular to the plate. Removal of In from the surface was performed by etching in concentrated HCl for several hours followed by thorough rinsing in ethanol. All samples obtained with this process showed no signs of In contamination. Powder X-ray patterns showed that samples crystallized in HoCoGa₅ structure without any additional peaks introduced by La alloying. In addition, magnetization measurements provided a more sensitive test of possible presence of magnetically ordered second phases. Both as grown and etched samples showed no sign of antiferromagnetic transition of CeIn₃. Electrical contacts were made with Epotek-H20E silver epoxy. In-plane resistivity was measured in Quantum Design MPMS and PPMS measurement systems from 0.35 to 300 K and in fields up to 90 kOe applied parallel and perpendicular to the c-axis. There is uncertainty in nominal resistivity values associated with sample geometry and uneven surfaces of etched samples. We measured several samples for each concentration in order to reduce measurement error which allowed us to estimate uncertainties in nominal values as well. The dimensions of the samples were measured by high precision optical microscope with $10\mu m$ resolution and average values are presented. Randomly chosen samples within each batch had no difference in their R(T) curves. Magnetization measurements were performed in MPMS-7 Quantum Design magnetometer in the magnetic field of 10kG, applied parallel and perpendicular to c axis.

III. RESULTS

The results of powder X-ray diffraction measurement taken at room temperature are summarized in Table 1 and shown in Fig. 1, together with the unit cell volume of LaCoIn₅. As expected, La doped samples have larger unit cell volume. The volume increase in the concentration range x=0-0.175 is consistent with expansion of the unit cell as La substitutes Ce in accordance with Vegard's law.

Fig. 2 shows the magnetic susceptibility for $Ce_{0.95}Nd_{0.05}CoIn_5$, $Ce_{0.85}La_{0.15}CoIn_5$, and $CeCoIn_5$,

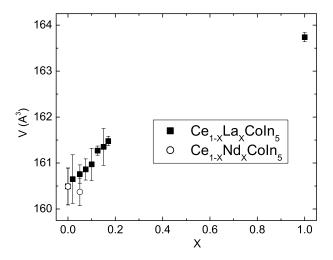


FIG. 1: Unit cell volume of $Ce_{1-x}La_xCoIn_5$ (x=0-0.175, 1) shown toghether with unit cell volume of $Ce_{0.95}Nd_{0.05}CoIn_5$

taken in the applied field of 10kOe. In the whole temperature range above T_c , the substitution of magnetic Ce³⁺ by nonmagnetic La³⁺ reduces susceptibility values in La doped sample when compared with undoped CeCoIn₅. Comparison of high temperature moments through Curie-Weiss analysis of the polycrystalline susceptibility average at high temperatures shows that approximately 14% Ce ions were substituted with La. No quantitative difference from undoped CeCoIn₅ was detected in high temperature susceptibility of 5% Nd doped sample. Low temperature magnetic susceptibility of Ce_{0.85}La_{0.15}CoIn₅ does not reveal any difference in Curie tail from pure material, thus ruling out Kondohole interpretation of La dilution (Fig. 2 inset). ¹² We also see broadening of the plateau-like feature in χ_c in Ce_{0.85}La_{0.15}CoIn₅ ascribed⁹ to thermal depopulation of Ce 4f levels. On the other hand, Nd impurities contribute to pronounced Curie tail at low temperatures. Subtraction of magnetic susceptibility of CeCoIn₅ from Ce_{0.95}Nd_{0.05}CoIn₅ in the normal state below 10K is consistent with approximately 8% of Nd³⁺ paramagnetic moment, result close to nominal stochiometric value and within rough approximation of our analysis.

Temperature dependent electrical resistivities normalized to their value at 300 K for $Ce_{1-x}La_xCoIn_5$ and $Ce_{0.95}Nd_{0.05}CoIn_5$ are presented in Fig. 3a. There are several key features to notice. Resistivities of all samples are weakly temperature dependant at high temperatures, and they pass through a maximum as temperature is decreased. This behavior is traditionally interpreted as a crossover from incoherent Kondo scattering to coherent Bloch states of heavy electrons in the Kondo lattice. In the case of $CeCoIn_5$ this drop, at least partially, could be attributed to depopulation of crystalline electric field levels. We observe decrease of T_{max} for higher La concentrations (Fig. 3a inset). At low temperatures, there is a clear suppression of T_c as more Ce ions are replaced by La (Fig. 3 inset). The increase of

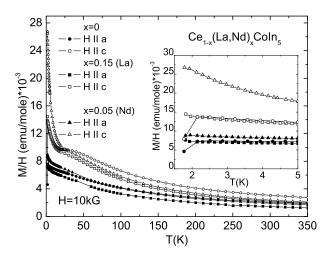


FIG. 2: Magnetic susceptibility of $Ce_{0.85}La_{0.15}CoIn_5$, $Ce_{0.95}Nd_{0.05}CoIn_5$ and $CeCoIn_5$. Low temperature susceptibility (inset) shows pronounced Curie tail with 5% of Nd substitution but no difference for 15% La substitution.

the normal state residual resistivity ρ_0 is probably due to disorder which contributes to increased conduction electron scattering. On the other hand, the resistive transition width sharpens with La alloying. It is interesting to note that $\text{Ce}_{1-x}\text{La}_x\text{CoIn}_5$ is not in the well defined Fermi liquid regime above T_c : the $\rho(T)$ curves above T_c do not show signs of T^2 dependence, as it has been reported for CeCu_2Si_2 . Depression of T_c in CeCoIn_5 seems to scale with ρ_0 values for both magnetic and nonmagnetic dopants, as seen by comparison of the $\rho(T)$ data of $\text{Ce}_{0.95}\text{Nd}_{0.05}\text{CoIn}_5$ and $\text{Ce}_{0.98}\text{La}_{0.02}\text{CoIn}_5$.

Fig. 4 shows the anisotropic upper critical field for $\text{Ce}_{1-x}\text{La}_x\text{CoIn}_5$, normalized to transition temperature in zero field for each value of x (values for x=0 were taken from previous report).¹⁴ The H_{c2} data were determined as a midpoint between onset and zero in resistivity from $\rho(T)$ curves at constant field and $\rho(H)$ curves at constant temperature. Adding La impurities results in depression of H_{C2} , however, anisotropy $\gamma = H_{c2}^a/H_{c2}^c$ remains at the same value of $\gamma \approx 2$ (inset in Fig. 4). Uncertainty in our estimate of γ decreases for higher field data, away from H=0 transition $(\text{T}/\text{T}_c \approx 1)$.

Assuming that Fermi surface properties of doped material do not change substantially in the dilute La limit, 15 it is reasonable to assume inverse proportionality between ρ and l, and therefore values of l_0 could be estimated from ρ_0 for the whole doping series $(l_0=\frac{A}{\rho_0})$ using the the value of constant A from reported l_0 and ρ_0 values for pure material. We obtain $l_0\approx 540\,\mbox{\normalfont\AA}$ for CeCoIn5 without La impurities. Fig. 5 shows the ratio of mean free path l_0 to in-plane superconducting coherence length ξ ($\xi^2(T)=\Phi_0/2\pi H_{c2}(T)$) for Ce_1_xLa_xCoIn5 obtained at T=T_c/2. In the whole doping range electronic system is in the clean limit which could explain nearly constant value of $\gamma=H_{c2}^a/H_{c2}^c$.

A comparison of the effects of La substitution on T_c

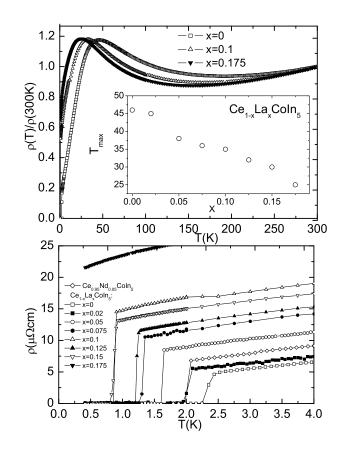


FIG. 3: (a) Electrical resistivity ρ normalized to its value at 300 K vs. temperature for $\text{Ce}_{1-x}\text{La}_x\text{CoIn}_5$ for $\text{x}=0,\,0.1$ and 0.175. T_{max} is shifted to lower temperatures with increased La substitution (inset) (b) Low temperature resistivity shows depression of T_c and increase in ρ_0 .

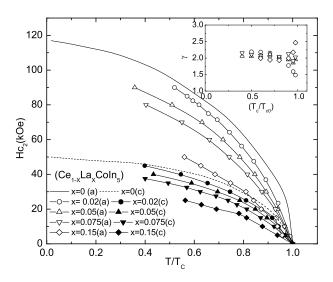


FIG. 4: Anisotropy in the upper critical field H_{c2} for $Ce_{1-x}La_xCoIn_5$ (x = 0-0.15). Inset shows value of $\gamma = H_{c2}^a/H_{c2}^c$ vs. $T_c/T_c(H=0)$ for varous La concentrations: x=0.02 (circles), x=0.05 (up triangles), x=0.075 (down triangles), x=0.15 (diamonds).

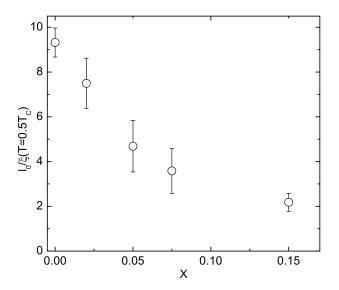


FIG. 5: Ratio of mean free path (l) to coherence length (ξ) for Ce_{1-x}La_xCoIn₅. Electronic system is in the clean limit already at T=T_c/2 for La concentrations x = 0-0.15

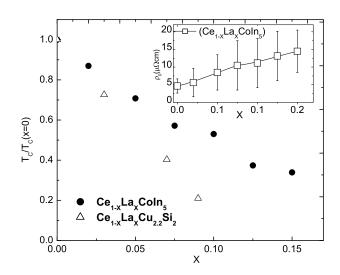


FIG. 6: Comparison of La doping on T_c of CeCoIn₅ (this work) and CeCu_{2.2}Si₂ (ref. 18). Inset shows increase in ρ_0 of Ce_{1-x}La_xCoIn₅ caused by La substitution.

in CeCoIn₅ and CeCu_{2.2}Si₂ is shown on Fig. 6.¹⁷ Doping results in depression of T_c in both cases but CeCoIn₅ is more robust to pair breaking arising from La impurities. The initial rate of T_c suppression is smaller than the rate seen in CeCu_{2.2}Si₂: [(0.056T_c)/(1% of La substitution) in CeCoIn₅ vs (0.085T_c)/(1% of La substitution in CeCu_{2.2}Si₂)]. La doping in CeCoIn₅ is associated with only modest increase in nominal residual resistivity values ρ_0 , shown in Fig. 6 inset. The ρ_0 values for x = 0 ($\sim 5\mu\Omega$ cm) in our experiment are in between values reported previously in literature (3.1 $\mu\Omega$ cm¹⁶ and $\sim 7\mu\Omega$ cm¹⁸).

IV. DISCUSSION AND CONCLUSIONS

The slope of H_{c2} vs T curve at T_c can be used to estimate zero temperature orbital critical field $H_{c2o}(0)$ using the weak - coupling formula for conventional superconductors in Werthamer-Helfand-Hohenberg model (WHH): $H_{c2o}(0) \approx 0.7 (H'_{c2}) T_c$. Table 1 shows estimates of H'_{c2} near T_c for doped samples, together with previously reported value for x = 0 for both crystalline directions.²⁰ All investigated samples have high initial slopes, as expected in the case of heavy fermion superconductors. ^{21,22} Values of $H_{c2o}(0)$ decrease with introduction of La impurities (Table 1). The paramagnetic limiting field $H_p(0) = \Delta_0/\mu_B\sqrt{2}$ (where Δ_0 is energy gap at T=0 and μ_B is Bohr magneton) for pure CeCoIn₅ $(T_c=2.3K)$ is well below the orbital critical field $H_{c2o}(0)$ for either s-wave $(\Delta_0 = 3.52k_BT_c)^{23}$ or d-wave pairing state $(\Delta_0 = 2.14k_BT_c)^{24}$ and our results indicate that this unusual situation is valid for investigated La doping range. We note that experimental values of upper critical field for $Ce_{1-x}La_xCoIn_5$ (x=0-0.15) samples are most likely below the values obtained by applying WHH model (Table 1), probably due to polarization of magnetic sublattice due to enhanced internal field along both crystalline axis.

It has recently been reported that T_c in CeCoIn₅ increases under applied pressure. 18 Negative chemical pressure should cause some decrease in T_c . In the lack of better approximation, we take bulk modulus of CeCoIn₅ to be the same as the one for CeIn₃ (650kbar),²⁵ and we calculate approximate chemical pressure ($P_{chemical}$) for each La concentration using $\frac{V_0 \partial P}{\partial V} \approx 650 \text{kbar}$. The results are shown in Table 1. Depression of T_c occurs at a rate $\frac{dT_c}{dP} \approx 0.43 \text{ K/kbar}$ - a slope that is an order of magnitude larger than reported increase of T_c under hydrostatic pressure. An order of magnitude difference from pure pressure effect on \mathcal{T}_c is likely to exceed error in estimation of bulk modulus, and therefore points to the conclusion that the pair breaking mechanisms which enter through disorder due to La alloving and increased scattering of Cooper pairs are dominant in CeCoIn₅. In contrast to conventional superconductors where nonmagnetic impurities have small effect on T_c , Cooper pairs formed in CeCoIn₅ are rather sensitive to La doping: 2% of La depresses T_c to the same value as $\sim 5\%$ of Nd.

The T_c suppression induced by the nonmagnetic La substitution in $Ce_{1-x}La_xCoIn_5$ is reminiscent of the pair breaking effect by magnetic impurities.²⁶ Although various factors may suppress T_c (an anisotropic scattering, for example),²⁷ we focus here on the scenario of $CeCoIn_5$ having an anisotropic gap $\Delta(\vec{k}_F)$ on the Fermi surface. This scenario is quite likely given the unconventional nature in many heavy-fermion materials.

It is known²⁸ that if Δ depends on the position at the Fermi surface, the critical temperature is suppressed by nonmagnetic scattering according to:

$$ln\frac{T_{c0}}{T_c} = \alpha \left[\psi(\frac{1+\mu}{2}) - \psi(\frac{1}{2}) \right], \mu = \frac{\hbar}{2\pi T_c}$$
 (1)

Here T_{c0} is the critical temperature of the material in the absence of all scattering, τ is the scattering time by nonmagnetic impurities, and $\alpha = 1 - \langle \Delta \rangle^2 / \langle \Delta^2 \rangle$ characterizes the gap anisotropy, $\langle ... \rangle$ stands for averaging over Fermi surface, and ψ is the digamma function. For a weak gap anisotropy, this result is due to Hohenberg, ²⁸ see also later publications. ^{29,30} It can be shown that in fact Eq. (1) holds for an arbitrary gap anisotropy. ³¹ For isotropic Δ , $\alpha = 0$, and we come to Anderson's theorem: $T_c = T_{c0}$. For pure d-wave order parameter, $\langle \Delta \rangle = 0$, and Eq. (1) describes the d-pair breaking by nonmagnetic scattering (which differs from the Abrikosov-Gor'kov result only by the factor of 2 in the definition of the parameter $\mu_m = \hbar/\pi T_c \tau_m$).

To analyze the $T_c(x)$ data shown in Fig. 6, one has to relate x to the scattering time τ , a nontrivial connection. We avoid this difficulty by assuming that the residual resistivity ρ_0 is proportional to $1/\tau$. Further, we exclude parameter T_{c0} from Eq. (1) by writing it for two values of x and subtracting the results:

$$ln\frac{T_2}{T_1} = \alpha[\psi(\frac{1+\mu_1}{2}) - \psi(\frac{1+\mu_2}{2})], \mu_{1,2} = \beta\frac{\rho_{1,2}}{T_{1,2}}$$
 (2)

where $T_{1,2} = T_c(x_{1,2})$ and β is a constant to be determined. Writing this equation for two different pairs $x_{1,2}$ one can determine the unknown α and β . This procedure yields values scattered around $\alpha = 0.5$ and $\beta = 0.2K/\mu\Omega cm$.

Hence, we find $\alpha = \langle \Delta \rangle^2 / \langle \Delta^2 \rangle \approx 0.5$ which implies a strongly anisotropic gap. Knowing the value of β we can estimate the scattering time using measured resistivities; for x=0 we obtain $\tau=\hbar/2\pi k_B\beta\rho\approx 1.3\times 10^{-12}s$. With the electronic specific heat coefficient $\gamma=290mJ/moleK^2$ we roughly estimate the Fermi velocity $v_F=\frac{\pi k_B}{e\sqrt{\gamma\tau\rho_0}}\approx 2\times 10^6 {\rm cm/s}$. This would correspond to the mean-free path $l\approx 260$ Å, a value smaller than expected but within factor of two of our determination of mean free path which is reasonable given the assumptions of average Fermi velocity and isotropic scattering.

In summary, diamagnetic pair breaking effect in CeCoIn₅ is consistent with picture of strongly anisotropic order parameter. Anisotropy in the upper critical field $\gamma = H_{c2}^a/H_{c2}^c$ does not change for x = (0-0.15) in Ce_{1-x}La_xCoIn₅, indicating electronic system in the clean limit

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x	$T_c(K)$	$a(\mathring{A})(\pm 0.007\mathring{A})$	$c(\mathring{A})(\pm 0.007\mathring{A})$	$V(\mathring{A})^3$	$-\frac{dH_{c2}}{dT}(kOe/K)$	$H_{c2o}(0)(kOe)$	$P_{chemical}(kbar)$
0	2.3	4.613	7.542	160.49 ± 0.4	$240(a),110\pm6(c)$	370(a), 170(c)	0
0.02	2.0	4.613	7.551	160.65 ± 0.53	$170\pm23(a),86\pm3(c)$	235(a), 119(c)	0.6
0.05	1.68	4.614	7.551	160.76 ± 0.2	$190\pm19(a),95\pm7(c)$	214(a), 107(c)	1.1
0.075	1.31	4.615	7.551	160.86 ± 0.23	$207\pm27(a),98\pm2(c)$	188(a), 89(c)	1.5
0.1	1.22	4.615	7.557	160.97 ± 0.35			2
0.125	0.86	4.623	7.546	161.27 ± 0.1			3.1
0.15	0.78	4.619	7.563	161.35 ± 0.4	$236\pm27(a),103\pm2(c)$	127(a), 55(c)	3.5
0.175	-	4.619	7.567	161.48 ± 0.1			
1.0	-	4.638	7.612	163.74 ± 0.1			

 160.37 ± 0.3

TABLE I: Properties of $Ce_{1-x}La_xCoIn_5$ doping series: T_c , lattice parameters, unit cell volumes, $H_{c2'}(T)$, calculated $H_{c2o}(0)$ from WHH model and approximate chemical pressure $P_{chemical}$ due to La alloying. Final row: properties of $Ce_{0.95}Nd_{0.05}CoIn_5$.

¹ Z. Fisk, J.L. Sarrao, J.L. Smith and J.D. Thompson Proc. Natl. Acad.Sci. USA **92**, 6663 (1995)

7.546

4.601

0.05(Nd) 2.0

- ² G. Bruls, D. Weber, B. Wolf, P. Thalmeir, B. Luthi, A. de Visser, A. Menovsky, Phys. Rev. Lett. 65, 2294 (1990)
- ³ B.S. Shivaram, J.J. Gannon, D. G. Hinks, Phys. Rev. Lett. 63, 1723 (1989)
- ⁴ K. Miyake, S. Schmitt-Rink and C. M. Varma, Phys. Rev B **34**, 6554 (1986)
- ⁵ H. Hegger, C. Petrovic, E. G. Moshopolou, M. F. Hundley, J. L. Sarrao, Z. Fisk and J. D. Thompson, Phys. Rev. Lett. 84, 4986 (2000)
- ⁶ C. Petrovic, R. Movshovich, M. Jaime, P. G. Pagliuso, M. F. Hundley, J. L. Sarrao, Z. Fisk and J. D. Thompson, Europhys. Lett. **53**, 354 (2001)
- ⁷ C. Petrovic, P. G. Pagliuso, M. F. Hundley, R. Movshovich, J. L. Sarrao, J. D. Thompson, Z. Fisk and P. Monthoux, J. Phys. Condens. Matter 13, L337 (2001)
- ⁸ Y. Kohori, Y. Yamato, Y. Iwamoto, T. Kohara, E. D. Bauer, M. B. Maple, J. L. Sarrao, Phys. Rev. B **64**, 134526 (2001)
- ⁹ N. J. Curro, B. Simovic, P. C. Hammel, P. G. Pagliuso, J. L. Sarrao, J. D. Thompson and G. B. Martins, Phys. Rev. B 64, 180514 (2001)
- ¹⁰ K. Izawa, H. Yamaguchi, Y. Matsuda, H. Shishido, R. Settai and Y. Onuki, Phys. Rev. Lett. 87, 057002 (2001)
- ¹¹ T. P. Murphy, Donovan Hall, E. C. Palm, S. W. Tozer, C. Petrovic, Z. Fisk, R. G. Goodrich, P. G. Pagliuso, J. L. Sarrao, and J. D. Thompson, Phys. Rev. B 65, 100514(R) (2002)
- J. M. Lawrence, T. Graf, M. F. Hundley, D. Mandrus, J. D. Thompson, A. Lacerda, M. S. Torikachvili, J. L. Sarrao and Z. Fisk, Phys. Rev B 53, 12559 (1996)
- ¹³ I. Sheikin, D. Braithwaite, J-P. Brison, W. Assmus and J. Floquet, J. Low Temp. Phys. **118**, 113 (2000)
- T. Muramatsu, N. Tateiwa, T. Kobayashi, K. Shimizu, K. Amaya, D. Aoki, H. Shishido, Y. Haga and Y. Onuki, J. Phys. Soc. Jpn. 70, 3362 (2001)

¹⁵ C.S. Jee, B. Andraka, J.S. Kim and G.R. Stewart, Phys. Rev B **43**, 2656 (1991)

-0.5

- ¹⁶ R. Movshovich, M. Jaime, J. D. Thompson, C. Petrovic, Z. Fisk, P. Pagliuso and J. L. Sarrao, Phys. Rev. Lett. 86, 5152 (2001)
- ¹⁷ U. Ahlheim, M. Winkelmann, P. van Aken, C. D. Bredl, F. Steglich and G.R. Stewart, J. Magn. Magn. Matter 76&77, 520 (1988)
- ¹⁸ M. Nicklas, R. Borth, E. Lengyel, P. G. Pagliuso, J. L. Sarrao, V. A. Sidorov, G. Sparn, F. Steglich and J. D. Thompson, J. Phys. Condens. Matter 13, L905 (2001)
- ¹⁹ N. R. Werthamer, E. Helfand and P. C. Hohenberg, Phys. Rev **147** 295 (1966)
- ²⁰ S. Ikeda, H. Shishido, M. Nakashima, R. Settai, D. Aoki, Y. Haga, H. Harima, Y. Aoki, T. Namiki, H. Sato and Y. Onuki, J. Phys. Soc. Jpn. **70**, 2248 (2001)
- ²¹ T. P. Orlando, E. J. McNiff, S. Foner and M. R. Beasley, Phys. Rev B **19**, 4545 (1979)
- ²² U. Rauchschwalbe, W. Lieke, C. D. Bredl, F. Steglich, J. Aarts, K. M. Martini and A. C. Mota, Phys. Rev. Lett 49, 1448 (1982)
- ²³ A. M. Clogston, Phys. Rev. Lett. **9**, 261 (1962)
- ²⁴ M.J. Graf, S-K. Yip, J.A. Sauls and D. Rainer, Phys. Rev. B **53**, 15147 (1996)
- ²⁵ G. Oomi, T. Kagayama and J. Sakurai, J. Mater. Process. Technol. 85, 220 (1999)
- ²⁶ A. A. Abrikosov and L. P. Gorkov, Soviet Physics JETP 12, 1243 (1961)
- ²⁷ J. Schmalian, private communication
- ²⁸ P. Hohenberg, Zh. Exp. Theor. Phys. **4**, 1208 (1963)
- ²⁹ D. Markowitz and L. P. Kadanoff, Phys. Rev. **131**, 563 (1963)
- ³⁰ A. I. Posazhennikova and M. B. Sadovski, JETP Lett. 63, 347 (1963)
- ³¹ V. G. Kogan, unpublished